PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6:	1	(11) International Publication Number: WO 98/27447
G02B 1/04, C08F 36/06	Aí	(43) International Publication Date: 25 June 1998 (25.06.98
(21) International Application Number: PCT/US (22) International Filing Date: 25 November 1997 ((30) Priority Data: 08/768,748 17 December 1996 (17.12.9 (70) Applicant: BAUSCH & LOMB INCORPORATED One Bausch & Lomb Place, Rochester, NY 14- (US). (72) Juventors: LAI, Yu-Chir. 23 Copper Woods, Pitts 14524 (US). BAMBURY, Ronald, E., 63 Matther Fairport, NY 14450 (US). (74) Agents: TROMAKS, John, E. et al.; Bausch & I corporated, One Bausch & Lomb Place, Roche 14604–2701 (US).	(25.11.9 (25.11.9 (US/US 604-270 ford, N ew Driv	Fi, GB, Hu, II., IP, KE, KG, KP, KR, MD, MK, MM, NN NO, NZ, PL, FT, RU, SE, SI, SK, TT, TT, TT, U, U, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW European patent (AT, BE, CH, DE, DK, ES, FI, FR, GE GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, B, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published With international search report.

(57) Abstract

Contact are made from polymerization product of a monomer mixture comprising a polybutadi-ene-based compound as represented by formula (1): E-B(AB) -E wherein: A

E-B-(AS)-E winteren: A is selected from the group consisting of a chemical bond, -OCO-, -COO-, -CONH-, -NHCO-, -OCOO-, -NHCOO- and -OCONH-; a is zero or an integer of at least 1; each B is independently represented by formula (II) where each of m, n, and p is independently or greater, provided that m + n + p is at least 10; R is hydrogen or C₁-C₄ alky; each E is independently a polymerizable terminal ethylenically unsaturated organic group.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	Prance	LU	Luxembourg	SN	Senegal .
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MĐ	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria.	HU	Hungary	ML	Meli	TT	Trinidad and Tobago
BJ	Benin	12	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Munitania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	zw	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand		
CM	Cameroou		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	8G	Singapore		

POLYBUTADIENE-BASED COMPOSITIONS FOR CONTACT LENSES

BACKGROUND OF THE INVENTION

Field of the Invention

This invention relates to novel contact lens materials, including hydrogel or rigid gas permeable (RGP) materials, which are made from the polymerization product of a monomer mixture including a polybutadiene-based compound endcapped with a polymerizable ethylenically unsaturated group.

Description of the Related Art

One class of materials useful for soft contact lenses is hydrogel polymers. A hydrogel is a hydrated, cross-linked polymeric system that contains water in an equilibrium state. Hydrogel contact lenses offer relatively high oxygen permeability as well as desirable biocompatibilty and comfort. Existing hydrogel soft contact lens materials are formed of copolymers based primarily on 2-hydroxyethylmethacrylate (Hema), N-vinyl-2-pyrrolidone (NVP) or other hydrophilic monomers, crosslinked with a crosslinking agent. Oxygen permeability of these materials is dependent on the water content of the hydrogel.

Another class of materials useful for soft contact lenses is elastomers. These materials are not hydrated, as are hydrogels, but are sufficiently flexible to be comfortable on the eve.

Contact lenses are also made from rigid, gas permeable (RGP) materials.

Generally, RGP materials are formed of copolymers based on silicon or fluorosilicon (meth)acrylates or itaconates, crosslinked with a crosslinking monomer. These materials are very rigid and are not hydrated, but due to the inclusion of a silicon or fluorosilicon monomer these materials have relatively high oxygen permeability.

It has been mentioned that contact lenses may be made of a polyolefin plastic, including polybutadiene. For example, U.S. Patent Nos. 5,290,548, 5,100,689, and 4,961,954 (Goldberg) describes methods for modifying the surface of contact lenses, and mentions that the lens may be made of a plastic such as polyolefins, in particular

polybutadienes. Additionally, it has been mentioned that butadiene may serve as a vinylic comonomer for hydrogel contact lenses, for example, US Patent No. 5,002,978 (Goldenberg). However, polybutadiene and butadiene do not readily copolymerize with conventional vinylic or (meth)acrylate monomers. Also the polymerization of polybutadiene and butadiene generally results in the loss of oxygen permeability.

The present invention provides a novel contact lens material based on polybutadiene compounds endcapped with a polymerizable ethylenically unsaturated group. These compounds can be readily combined with conventional contact lens monomers, so as to obtain materials in which the desired properties, such as hardness, oxygen permeability, and water content are readily controlled by adjusting monomer concentration. Additionally, these compounds can be molded into lenses by conventional curing processes using free radical polymerization.

SUMMARY OF THE INVENTION

The invention relates to contact lenses made from the polymerization product of a monomer mixture comprising a polybutadiene-based compound as represented by formula (I)

wherein:

A is selected from the group consisting of a chemical bond, -OCO-, -COO-, -CONH-, -NHCO-, -OCOO-, -NHCOO- and -OCONH-;

a is zero or an integer of at least 1;

each B is independently represented by the formula (II)

where each of m, n, and p is independently 0 or greater, provided that m + n + p is at least 10:

R is hydrogen or C1 - C4 alkyl;

each E is independently a polymerizable terminal ethylenically unsaturated organic group represented by the formula (III)

wherein R2 is hydrogen or methyl;

 R^3 is selected from the group consisting of hydrogen, alkyl having 1 to 6 carbon atoms, and a $-CO-Y^*-R^1$ radical wherein Y^* is -O-, -S-, or -NH- and R^1 is a C_1 to C_{12} alkyl;

each of X and R^4 is independently selected from the group consisting of -OCO-, -COO-, -CONH-, -NHCO-, -OCOO-, -NHCOO- and -OCONH-;

Ar is an aromatic radical having 6 to 30 carbon atoms;

each of t and y is independently 0 or an integer of 1 to 6; and each of u, y and w is independently 0 or 1.

A preferred class of compounds of Formula I include those represented by the formula

wherein E is a group of the formula

$$\begin{array}{c} R^2 \\ CH_2 = C - COO - (CH_2)_{\widetilde{y}} - (NHCOO)_{V} \end{array}$$

and B is

wherein R2 is hydrogen or methyl;

y is 0 or an integer of 1 to 6;

v is 0 or 1; and

m + n is at least 10.

DETAILED DESCRIPTION OF THE INVENTION

The contact lens materials of the invention are formed from the polymerization product of a monomer mixture comprising a polybutadiene-based compound as represented by formula (I):

wherein:

A is selected from the group consisting of a chemical bond, -OCO-, -COO-, -CONH-, -NHCO-, -OCOO-, -NHCOO- and -OCONH-;

a is zero or an integer of at least 1;

B is represented by the formula (II)

wherein m, n, and p is independently 0 or greater, provided that m + n + p is at least 10;

R is hydrogen or C1 - C4 alkyl;

each E is independently a polymerizable terminal ethylenically unsaturated organic group represented by the formula (III)

wherein R2 is hydrogen or methyl;

 R^3 is selected from the group consisting of hydrogen, an alkyl having 1 to 6 carbon atoms, and a $-CO-Y'-R^1$ radical wherein Y' is -O-, -S-, or -NH- and R^1 is C_1 to C_{12} alkyl;

X and R 4 are independently selected from the group consisting of -OCO-, -COO-, -CONH-, -NHCO-, -OCOO-, -NHCOO- and -OCONH-;

Ar is an aromatic radical having 6 to 30 carbon atoms; each of t and y is independently 0 or an integer of 1 to 6; and each of u. y and w is independently 0 or 1.

One preferred class of compounds of Formula I include those represented by the formula:

i.e., compounds of formula (I) wherein a is one and A is a chemical bond.

The polybutadienes of formula (I) specifically include compounds based on 1,3 butadiene (where R is hydrogen) and 2-methyl-1,3 butadiene (where R is methyl).

Additionally, preferred B radicals include:

where m + n is at least 10.

A preferred class of E radicals have the formula:

$$R^2$$

CH₂=C-COO-(CH₂)_V-(NHCOO)_V

where R2, y and v are as defined for formula (I).

Various formula (I) compounds are commercially available. Examples are the methacrylate-endcapped polybutadiene (molecular weight 11,000) available from Aldrich (Milwaukee, WI), and the acrylate-capped polybutadiene (molecular weight 3,000) available from AtoChem (Philadelphia, PA).

Other compounds can be prepared by methods known in the art. As an example, the polybutadiene-based compounds of formula (I) may be prepared by endcapping the polybutadienes of formula (IV):

where R⁵ is a reactive group containing hydroxyl or amino. More specifically, compounds of formula (IV) can be reacted with ethylenically unsaturated radicals according to conventional methods.

As a first example, polybutadiene-based compounds of formula (IV) containing terminal hydroxyl functionality may be reacted with isocyanatoethylmethacrylate to form compounds of Formula (I) wherein E is

Alternatively, polybutadiene-based compounds of formula (IV) containing terminal hydroxyl functionality may be reacted with (meth)acryloyl chloride to provide a (meth)acrylate terminal radical, or with vinyl chloroformate to provide a vinyl carbonate terminal radical. Various hydroxy-terminated polybutadiene-based compounds of formula (IV) are commercially available from Nippon Soda Co. (Japan).

Polybutadiene-based compounds of formula (IV) containing terminal amino functionality may be reacted with (meth)acryloyl chloride to provide a terminal (meth)acrylamide terminal radical, or with vinyl chloroformate to provide a vinyl carbamate end group.

Monomer mixtures comprising the ethylenically unsaturated monomers of formula (I) may be polymerized by free radical polymerization to form contact lenses according to conventional methods. It has been found that such polymeric shaped articles have sufficiently high oxygen permeability, clarity and strength for use as contact lens materials.

The monomer mixture may employ a compound of formula (I) as the sole monomeric component. Alternately, the monomer mixture may comprise, in addition to the formula (I) compound, conventional monomers used for contact lens applications.

As one example, compounds of formula (I) may be copolymerized with at least one hydrophilic monomer to form a hydrophilic, optically clear copolymer useful as a soft, hydrogel contact lens material. Suitable hydrophilic monomers include: unsaturated carboxylic acids, such as methacrylic and acrylic acids; (meth)acrylic substituted alcohols. such as 2-hydroxyethyl methacrylate and 2-hydroxyethyl acrylate; glyceryl methacrylate; vinyl lactams, such as N-vinyl-2-pyrrolidone; and acrylamides, such as methacrylamide, N.N-dimethylacrylamide, N-(2-hydroxyethyl)-methacrylamide, and N-methacryloyl glycine. Still further examples are the hydrophilic vinyl carbonate or vinyl carbamate monomers disclosed in U.S. Pat. No.5,070,215, and the hydrophilic oxazolone monomers disclosed in U.S. Pat. No.4,910.277. Preferably, the polybutadiene-based compound is included in the initial monomeric mixture at about 50 to about 80 percent by weight, and at least one hydrophilic monomer is included at about 20 to about 50 percent by weight. Either the polybutadiene-based compound or the hydrophilic monomer may function as a crosslinking agent (a crosslinker being defined as a material having multiple polymerizable functionalities). Optionally, a separate crosslinker may be employed in the initial monomeric mixture to provide a crosslinked polymeric article, generally at 0.1 to 20 percent by weight when present.

Additionally, the macromonomers may be copolymerized with monomers such as methylmethacrylate, an itaconate ester, or fluorinated derivatives thereof to form a RGP contact lens materials. Generally, the monomer mixture will further include one of the forementioned hydrophilic monomers as a wetting agent, and optionally a crosslinking agent.

The monomeric mixtures including the formula (I) compound may be polymerized by free radical polymerization, usually in the presence of heat or ultraviolet irradiation. Minor amounts of a free-radical initiator may be included in the monomeric mixture, generally at about 0.1 to about 5 percent by weight. In producing contact lenses, the initial monomeric mixture may be cured in tubes to provide rod-shaped articles, which are

then cut into buttons. The buttons may then be lathed into contact lenses. Alternately, contact lenses may be cast directly in molds from the monomeric mixtures, such as by spincasting and static casting methods. Spincasting methods are disclosed in U.S. Pat. Nos. 3,408,429 and 3,660,545, and static casting methods are disclosed in U.S. Pat. Nos. 4,113,224, 4,197,266, and 5,271,875. As an additional method, U.S. Pat. No. 4,555,732 discloses a process where an excess of a monomeric mixture is cured by spincasting in a mold to form a shaped article having an anterior lens surface and a relatively large thickness, and the posterior surface of the cured spincast article is subsequently lathed to provide a contact lens having the desired thickness and posterior lens surface.

The following examples illustrate various preferred embodiments of the present invention

EXAMPLES 1 and 2

Two monomer mixtures were prepared. The first mixture contained: a methacrylate-endcapped polybutadiene obtained from Aldrich (Milwaukee, WI) having a molecular weight of 11,000 (60 parts by weight); toluene (40 parts by weight); and a UV initiator, DAROCUR 1173 (0.5 parts by weight). The second mixture contained: an acrylate-endcapped polybutadiene obtained from AtoChem (Philiadelphia, PA) having a molecular weight 3,000 (60 parts by weight); toluene (40 parts by weight); and DAROCUR 1173 initiator (0.5 parts by weight).

Two series of films were cast from the two mixtures between glass plates by subjecting the mixtures to ultraviolet irradiation for about two hours. The films were separated from the glass plates, extracted in alcohol, dried and re-extracted with boiling water. The extracted films were then placed in phosphate-buffered saline.

The resultant films were optically clear. Oxygen permeability was measured by the "single-chamber method" (Fatt, I; Rasson, JE; Melpolder, JB, <u>International Contact Lens Clinic</u>, vol. 14(10), p 389 1987). Using this method, oxygen permeabilities are measured in units of Dk (Dk equals 10⁻¹¹ cm² mL O₂/s mL mmHg). Oxygen permeability of the first set of films (based on the methacrylate-capped polybutadiene) averaged 50 Dk.

and oxygen permeability of the second set of films (based on the acrylate-capped polybutadiene) average 33 Dk,

EXAMPLES 3-10

A series of monomer mixtures was prepared by mixing the methacrylate-capped and acrylate-capped polybutadiene described in Example 1 and 2 with N,N-dimethyl acrylamide (DMA) at various ratios. (Examples 3-6 in Table 1 are monomer mixtures including the methacrylate end-capped polybutadiene (MW 11,000), and Examples 7-10 are monomer mixtures including the acrylate end-capped polybutadiene (MW 3,000). The monomer mixtures further included 30.0 weight % toluene and 0.5 weight % DAROCUR 1173 initiator. The monomer mixtures were then cast into films as described in Examples 1 and 2 and extracted as described in Examples 1 and 2. Oxygen permeability of the resultant films was measured as in Examples 1 and 2. Water content of the hydrated films was measured gravimetrically. Mechanical testing of the films was conducted in buffered saline on an Instron instrument, according to a modified ASTM method D-1708 (tensile properties) and D-1938 (tear strength) procedures. Alcohol extractables were determined gravimetrically. The properties of the films are listed in Table I.

TABLE 1

Poly(butadiene) Compound		MW 11,000				MW 3,000		
Examples:	3	4	5	6	7	8	9	10
% End-capped polybutadiene	80	70	60	50	80	70	60	50
% DMA	20	30	40	50	20	30	40	50
Properties					 			
					1			
% Extr.	8.6	11.2	11.7	15.5	4.1	6.7	7.4	8.8
% water	9.7	20.0	31.6	43.1	4.1	18.6	29.4	43.0
Dk	30.6	25.5	25.9	27.2	24.6	23.6	23.5	26.5
Modulus g/mm2	23	22	20	15	41	36	31	24
Tear g/mm	7.5	6.3	5.6	4.2	5.3	4.8	3.7	3.2

The mechanical properties (modulus and tear strength) were acceptable for contact lens applications.

For comparison purposes, a hydrated hydrogel of polymacon (a commercial hydrogel containing HEMA as the primary hydrophilic monomer) has a water content of about 38 weight % and an oxygen permeability of about 9 Dk. In contrast, the materials of the invention exhibited a markedly higher oxygen permeability, even the materials having water contents much lower than 38 weight %.

We claim:

 A contact lens formed from the polymerization product of a monomer mixture comprising a polybutadiene-based compound having the following structure

E-B(AB).-E

in which

each A is independently selected from the group consisting of a chemical bond or -OCO-, -COO-, -CONH-, -NHCO-, -OCOO-, -NHCOO- and -OCONH-; a is zero or an integer of at least 1;

each B is independently represented by the formula

where each of m, n, and p is independently 0 or greater, provided such that m + n + p is at least 10:

R is hydrogen or C1 - C4 alkyl;

each E is independently a polymerizable terminal ethylenically unsaturated organic group having the formula:

$$\begin{array}{ccc} & H & R^2 \\ I & I & I \\ R^3 - C = C - (CH_2)_{I}(X)_{L}(Ar)_{W}(CH_2)_{V}(R^4)_{V} \end{array}$$

wherein R2 is hydrogen or methyl;

 R^3 is selected from the group consisting of hydrogen, an alkyl having 1 to 6 carbon atoms, and a $-CO-Y'-R^1$ radical wherein Y' is -O-, -S-, or -NH- and R^1 is a C_1 to C_{12} alkyl;

each of X and R⁴ is independently selected from the group consisting of -OCO -,
-COO -, -CONH -, -NHCO -, -OCOO -, -NHCOO - and -OCONH -;
Ar is an aromatic radical having 6 to 30 carbon atoms;
each of t and y is independently 0 or an integer of 1 to 6;
and each of u, v and w is independently 0 or 1.

 The contact lens of claim 1 wherein the monomer mixture further comprises a hydrophilic monomer.

- The contact lens of claim 2 wherein the monomer mixture contains about
 to about 80 weight percent of the polybutadiene-based compound of claim 1.
- 4. The contact lens of claim 2 wherein the monomer mixture contains about 60 to about 70 weight percent of the polybutadiene-based compound of claim 1.
- 5. The contact lens of claim 2 wherein the hydrophilic monomer includes at least one member selected from the group consisting of N,N- dimethylacrylamide, N-vinyl-2-pyrrolidone, 2-hydroxyethyl methacrylate, N-(2-hydroxyethyl)-methacrylamide, glyceryl methacrylate and methacrylic acid.
- The contact lens of claim 2 wherein the monomer mixture further comprises a polymerization initiator.
- The contact lens of claim 2 wherein the monomer mixture further comprises a crosslinking agent.
 - 8. The contact lens of claim 1 wherein each E is

$$R^2$$
 $CH_2 = C - COO - (CH_2)_y - (NHCOO)_y$

wherein R2 is hydrogen or methyl;

y is 0 or an integer of 1 to 6; and v is 0 or 1.

9. The contact lens of claim 8 wherein each E is

 The contact lens of claim 9, wherein the polybutadiene-based compound has the formula

and a molecular weight of 2,000 to 15,000.

 A hydrogel that is the hydrated polymerization product of a monomer mixture comprising the polybutadiene-based compound having the following structure E-B(AB)_E

in which

each A is independently selected from the group consisting of a chemical bond or -OCO -, -COO -, -CONH -, -NHCO -, -OCOO -, -NHCOO - and -OCONH -:

a is zero or an integer of at least 1;

each B is independently represented by the formula

where each of m, n, and p is independently 0 or greater, provided such that m + n + p is at least 10:

R is hydrogen or C1 - C4 alkyl;

each E is independently a polymerizable terminal ethylenically unsaturated organic group having the formula:

wherein R2 is hydrogen or methyl;

 R^3 is selected from the group consisting of hydrogen, an alkyl having 1 to 6 carbon atoms, and a $-CO-Y'-R^1$ radical wherein Y' is -O-, -S-, or -NH- and R^1 is a C1 to C12 alkyl:

each of X and R⁴ is independently selected from the group consisting of - OCO -, - COOH -, - NHCO -, - OCOO -, - NHCOO - and - OCONH -;

Ar is an aromatic radical having 6 to 30 carbon atoms; each of t and y is independently 0 or an integer of 1 to 6; and each of u,v and w is independently 0 or 1.

12. The hydrogel of claim 11 wherein the monomer mixture further comprises at least one hydrophilic monomer.

A. CLASSIFICATION OF SUBJECT MATTER						
G 02 B 1/04,C 08 F 36/06						
			1			
According to	International Patent Classification (IPC) or to both national classifi	cation and EBC6	1			
	SRARCHED					
	ocumentation searched (classification system followed by classification	on symbols)				
	12 B,C 08 F,C 08 J		1			
	2 2 0 00 1 70 00 0		1			
D						
Document	ion searched other than minimum documentation to the extent that si	ich documents are included in the licids st	archeo			
Electronic d	ata base consulted during the international search (name of data base	and, where practical, search terms used)				
		•	1			
			1			
C. DOCUM	IENTS CONSIDERED TO BE RELEVANT					
Category*	Citation of document, with indication, where appropriate, of the rel	evant passages	Relevant to claim No.			
A	US 5290548 A		1,3,4			
	(GOLDBERG et al.) 01	March	1			
	1994 (01.03.94),		ł			
	claims 1,10-12	*4 **)	1			
	(cited in the applica	CION).	į			
A	US 5100689 A		1,3,4			
••	(GOLDBERG et al.) 31	March	1,5,5			
	1992 (31.03.92),					
	claims 1,6,12, column 7,					
	lines 20-35					
	(cited in the applica	tion).				
A	US 4961954 A		1,3,4			
1	(GOLDBERG et al.) 09	October	1			
	1990 (09.10.90),	14				
	claims 1,6, column 5, 61 - column 6, line 3	Tine				
1	(cited in the applica					
1	(creed in the apprica					
	L					
X Fur	ther documents are listed in the continuation of box C.	Patent family members are listed	in annex.			
* Special co	stegories of cited documents :	T later document published after the in	ternational filing date			
A document defining the seneral case of the art which is not or priority date and not in conflict with the application but						
conte	sered to be of particular relevance	invention				
"E" earlier document but published on or after the international filing date "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to						
"L' document which may throw doubts on priority claim(s) or involve an inventive step when the document is taken alone						
account or other special reason (as specified) earned the considered to involve an inventive step when the						
O document referring to an oral discloture, use, exhibition or document is combined with one or more other such document of ments, such combination being obvious to a person stalled						
P' document published more to the international filing date but in the art.						
later than the priority date claimed & document member of the same patent family						
Date of the	Date of the actual completion of the international search 19 February 1998 10 no no no					
1	epragry 1990	1 0. 03. 98				
Name and	mailing address of the ISA	Authorized officer				
	European Patent Office, P.B. 5818 Patentiaan 2 NL - 2280 HV Rijswijk					
l	Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.	HAUSWIRTH e.h.				
(Fax (+31-70) 340-3016					

-2-

INTERNATIONAL SEARCH REPORT Internation optication No PCT/US 97/22060

	-2-	PCT/US 91/22060						
(Continue	inuation) DOCUMENTS CONSIDERED TO BE RELEVANT y * Citagon of document, with indication, where appropriate, of the relevant passages Relevant to claim No.							
_avegory -	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.						
A	US 5002978 A	1,11						
	· ·							

ANHANG

zum internationalen Recherchen-bericht über die internationale Patentanmeldung Nr.

In Recherchenhamicht

ANNEX

to the International Search Report to the International Patent Application No.

ANNEXE

au rapport de recherche inter-national relatif à la demande de brevet international n°

PCT/US 97/22060 SAE 178542

Datus der

Mitalianian) dar

In diesea Anhang sind die Mitglieder der Patentalien der in debenger der Patentalien der in der Patentalien de

Datum doe

angefü Pat in Docum dans 1	Recherchenbericht hrtes Patentdokument ent document cited i search report ent de brevet cité e rapport de recherche	Datum der Veröffentlichung Publication date Date de publication	Mitolied(er) der Patentfamilie Patent family member(s) Membre(s) de la familie de brevets	Datum der Veröffentlichung Publication date Date de publication	
US A	A 529054B	01-03-94	1738/34/19/19/19/19/19/19/19/19/19/19/19/19/19/	288223802665026652523425652778446555547775555-161449	
US	A 5100689	31-03-92	120034447333410040000000000000000000000000		

•					
*.				50 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	03-04-9-2
	US Ā	4961954	09-10-90	\$\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	28007788007695767447573347577777877798457678757911-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1
	US A	5002978	26-03-91 -	AT E 1 19797 AT	15-03-95 15-07-90 15-07-90 15-03-95 15-03-